

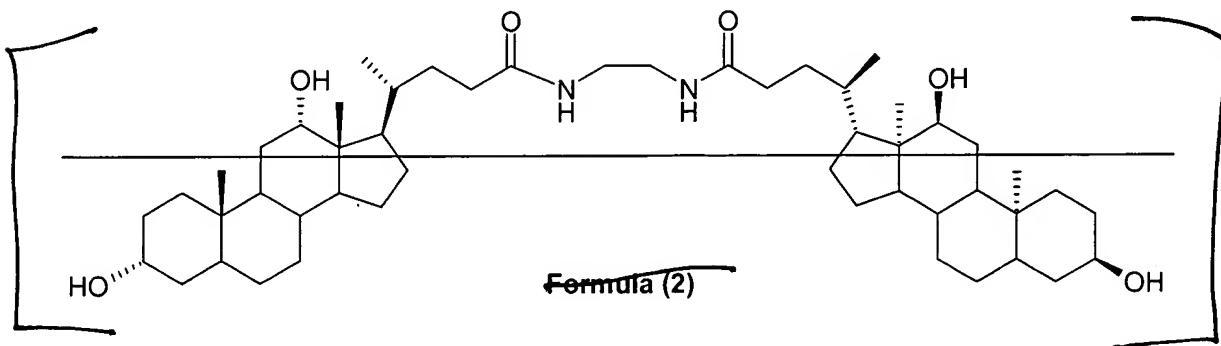
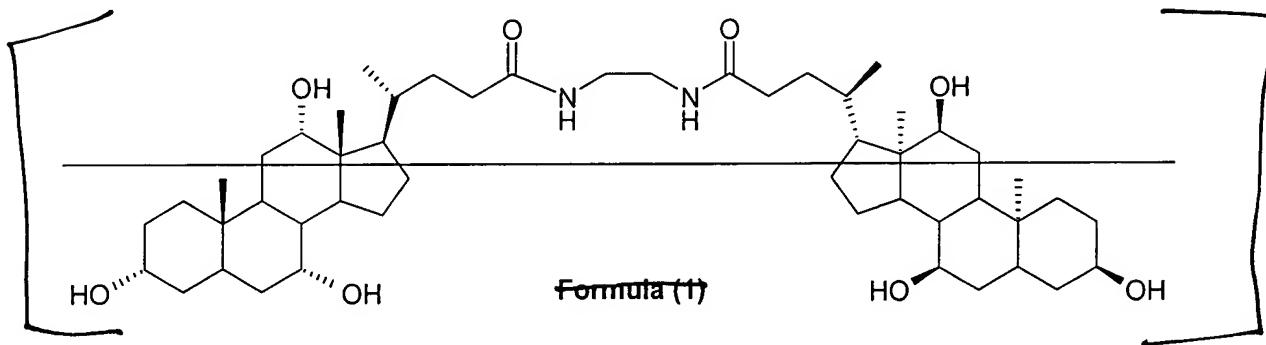
**AMENDMENTS TO THE CLAIMS**

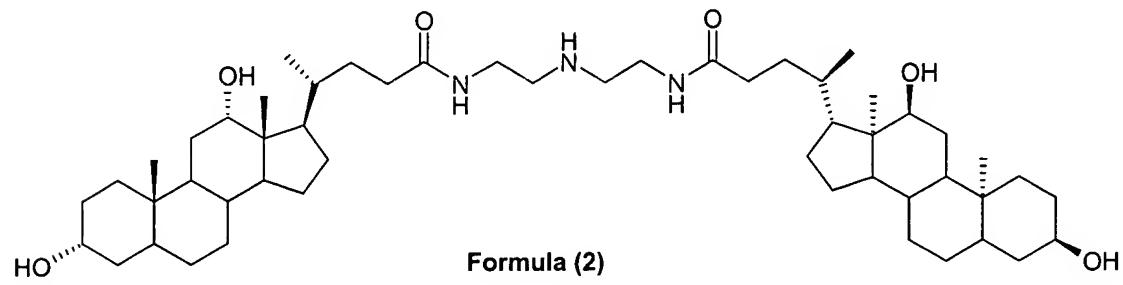
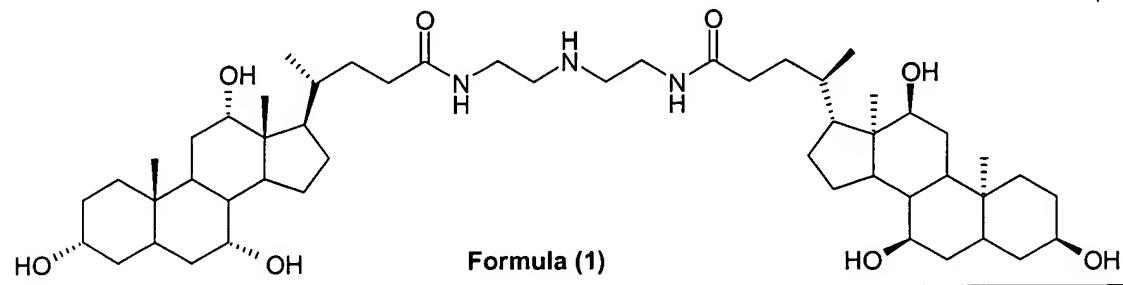
This listing of claims will replace all prior versions and listings of claims in the application:

**LISTING OF CLAIMS**

We Claim

1. (currently amended) Antifungal steroidal dimers,  $N^1, N^3$ - diethylenetriamine bis [cholic acid amide] of formula (1), and  $N^1, N^3$ - diethylenetriamine bis [deoxycholic acid amide] of formula (2)





comprising amphiphilic topology as shown below:

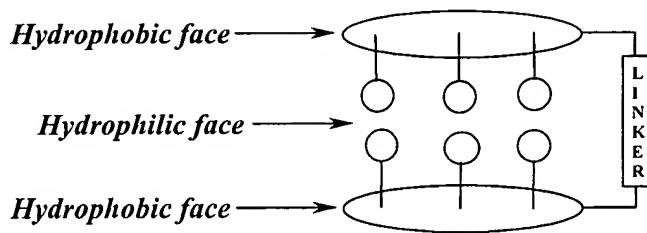
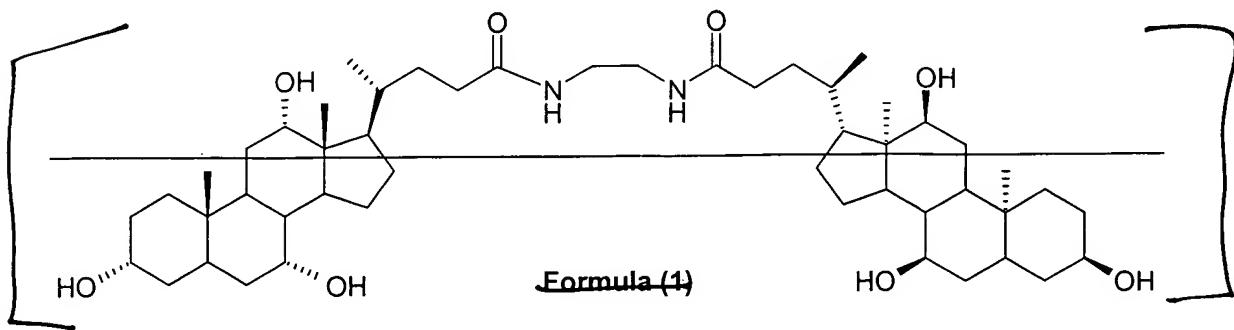


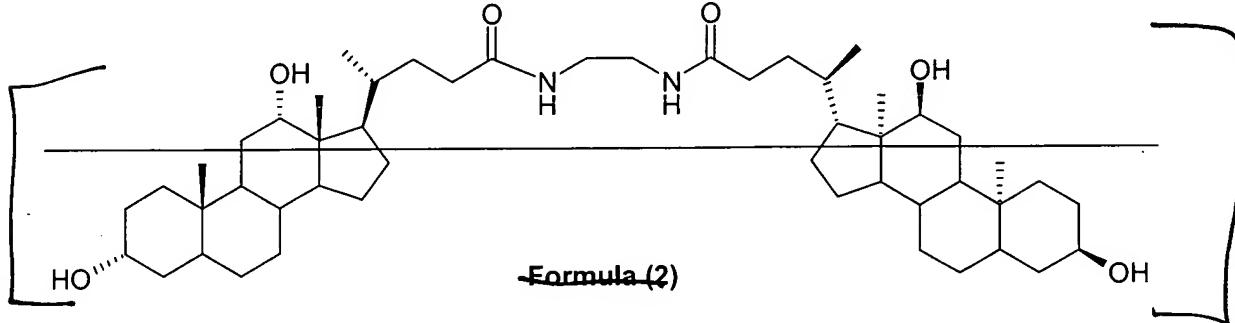
Figure (3)

2-6. (canceled)

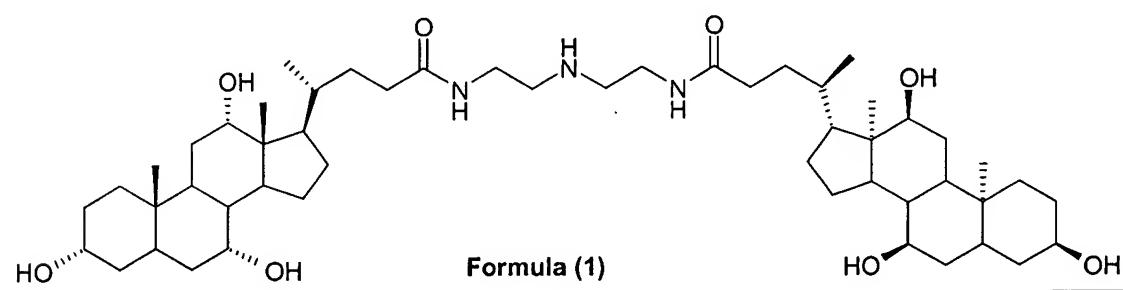
7. (currently amended) A process method for the preparation of steroidal dimers N<sup>1</sup>, N<sup>3</sup>-diethylenetriamine bis [cholic acid amide] and N<sup>1</sup>, N<sup>3</sup>-diethylenetriamine bis [deoxycholic acid amide] having structural formula (1) and (2) respectively,



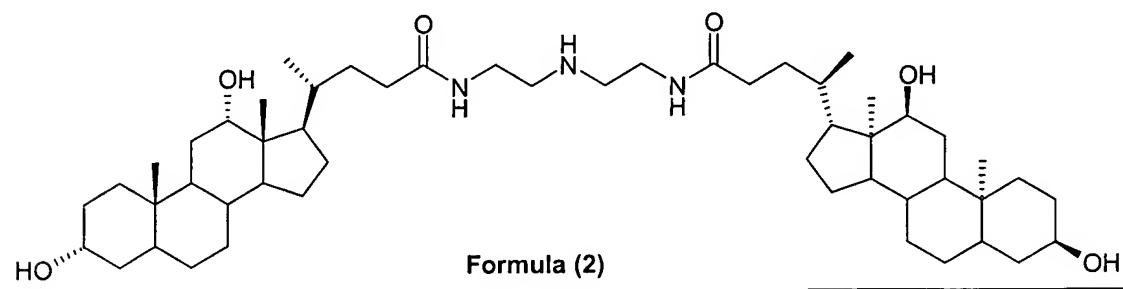
**Formula (1)**



**Formula (2)**



**Formula (1)**



**Formula (2)**

comprising amphiphilic topology as shown below:

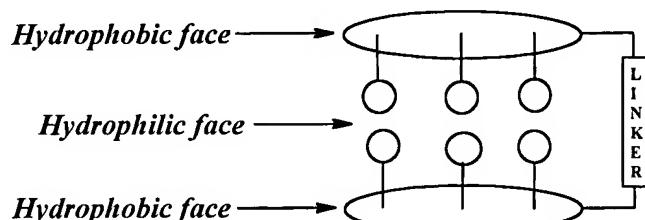


Figure (3)

said process method comprising steps of,

- a. preparing a solution of N-succinimidyl ester of bile acids in an organic solvent at a temperature ranging between 10 to 50 °C;
- b. adding diethylenetriamine to the solution of step (a) followed by stirring the same for a time duration ranging from 1 to 5 h at a temperature ranging between 20 to 70 °C to obtain a reaction mixture;
- c. quenching the reaction mixture of step (b) with ice to a form containing crude products having structural formula (a) (1) and (b) (2), and
- a. separating the crude products of step (c) and purifying the same to obtain the compound of formula (1) or (2).

8. (canceled)

9. (currently amended) A process as claimed in The method of claim 7, wherein the organic solvent[s] is selected from a group comprising chlorinated solvents such as chloroform and dichloromethane or polar aprotic solvents such as dimethylformamide and dimethylsulfoxide.

10-13. (canceled)